

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



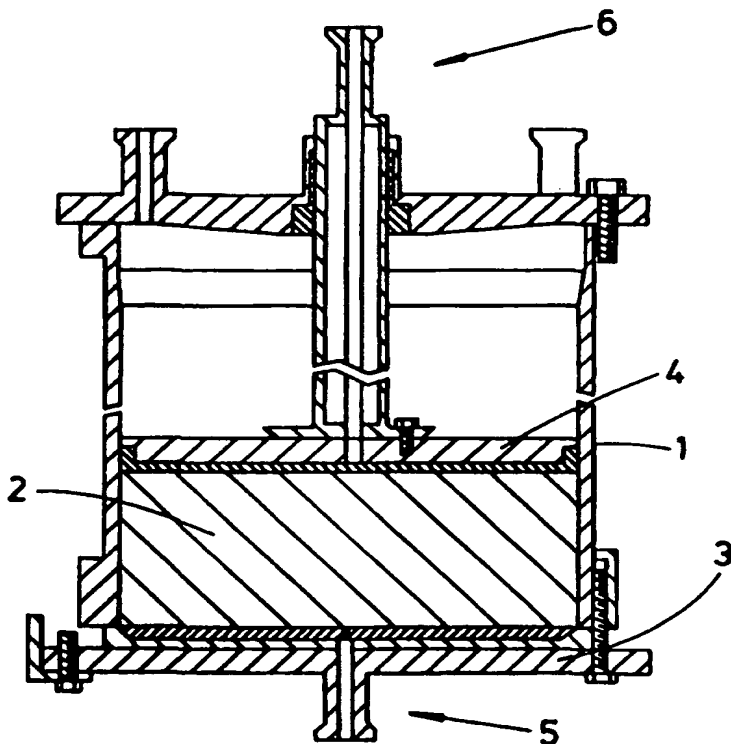
INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : G01N 30/60, B01D 15/08		A1	(11) International Publication Number: WO 96/26436
			(43) International Publication Date: 29 August 1996 (29.08.96)
(21) International Application Number: PCT/SE96/00206		(81) Designated States: AU, CA, JP, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).	
(22) International Filing Date: 16 February 1996 (16.02.96)			
(30) Priority Data: 9500635-9 21 February 1995 (21.02.95) SE		Published With international search report. In English translation (filed in Swedish).	
(71) Applicant (for all designated States except US): PHARMACIA BIOTECH AB [SE/SE]; S-751 82 Uppsala (SE).			
(72) Inventor; and (75) Inventor/Applicant (for US only): PETTERSSON, Conny [SE/SE]; Enspännargatan 33, S-165 57 Hasselby (SE).			
(74) Agents: BERGANDER, Håkan et al.; Pharmacia & Upjohn, Patent Dept., S-751 82 Uppsala (SE).			

(54) Title: COLUMN FOR CHROMATOGRAPHY

(57) Abstract

The present invention relates to a liquid-chromatography column constructed from a column tube (1) containing a chromatographic matrix (2), a fluid inlet (3), a fluid outlet (4) and a respective adapter (5 and 6) placed at the inlet and the outlet respectively. The column is characterized in that respective adapters (5 and 6) include an end-plate (7) and a perforated plate (8), such as to form a gap (9) between the end-plate and the perforated plate (8). The inventive column is particularly suited for large-scale chromatography in which matrices of small particle sizes are used.



FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AM	Armenia	GB	United Kingdom	MW	Malawi
AT	Austria	GE	Georgia	MX	Mexico
AU	Australia	GN	Guinea	NE	Niger
BB	Barbados	GR	Greece	NL	Netherlands
BE	Belgium	HU	Hungary	NO	Norway
BF	Burkina Faso	IE	Ireland	NZ	New Zealand
BG	Bulgaria	IT	Italy	PL	Poland
BJ	Benin	JP	Japan	PT	Portugal
BR	Brazil	KE	Kenya	RO	Romania
BY	Belarus	KG	Kyrgyzstan	RU	Russian Federation
CA	Canada	KP	Democratic People's Republic of Korea	SD	Sudan
CF	Central African Republic	KR	Republic of Korea	SE	Sweden
CG	Congo	KZ	Kazakhstan	SG	Singapore
CH	Switzerland	LI	Liechtenstein	SI	Slovenia
CI	Côte d'Ivoire	LK	Sri Lanka	SK	Slovakia
CM	Cameroon	LR	Liberia	SN	Senegal
CN	China	LT	Lithuania	SZ	Swaziland
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	LV	Latvia	TG	Togo
DE	Germany	MC	Monaco	TJ	Tajikistan
DK	Denmark	MD	Republic of Moldova	TT	Trinidad and Tobago
EE	Estonia	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	UG	Uganda
FI	Finland	MN	Mongolia	US	United States of America
FR	France	MR	Mauritania	UZ	Uzbekistan
GA	Gabon			VN	Viet Nam

Column for Chromatography

Technical field

The present invention relates to a column for liquid chromatography.

Description of known techniques

When practicing liquid chromatography on a porous matrix, a liquid containing a dissolved compound is allowed to pass through the matrix, wherewith the compound flows through the matrix while passing through one or more adsorption/desorption stages.

Matrices intended for liquid chromatography are normally comprised of particles, e.g. beads, that are packed together in a column tube to form a bed. The bed is normally held in place in the tubular column with the aid of two adapters, each covering a respective end of the bed and therewith also the cross-sectional area of the column. One of the adapters will often include an inlet for elution agent which prior to penetrating the matrix bed passes through a perforated plate which distributes the flow uniformly over the end area of the bed. The other adapter has an outlet for elution agent, which prior to entering the outlet also passes through a perforated plate which gathers the flow uniformly across the end area of the bed prior to the elution agent exiting through the outlet. It is normal to place a very fine net between the perforated plate and the gel bed, to prevent the ingress of gel material into respective adapters. Examples of perforated plates are filter plates, gratings, coarse nets, apertured plates and discs.

With the intention of minimizing diffusion and zone spreading during chromatography, the perforated plate, either with or without a fine net, is placed tightly against the inlet area of the bed at respective ends of the column. This is done to obviate the risk of particles swirling up from the bed.

According to known technique, perforated plates are placed against the gel bed and immediately adjacent the end-plate of respective adapters.

The problems of known techniques

Conventional perforated plates, such as gratings, etc., have been found to present problems in large scale chromatographic processes in which the gel matrix consists of small beads, which require greater bed packing pressures and are operated at high rates of flow. By large-scale columns is meant here column tubes that have a diameter greater than or equal to 50 mm. A test which functions well on a small scale will often present problems in large-scale chromatography performed under the aforesaid conditions.

More specifically, the earlier known perforated plates impede the radial flow of eluting solution and sample solution respectively in the end-areas of the column, giving rise to an uneven plug flow. This uneven plug flow will often result in a more rapid flow in the middle of the column and slower flow out towards the periphery of the gel-bed cross-section. This is shown in the chromatogram by broad peaks with high degrees of dilution. The plate number of the column will therewith be lowered. Because of the high packing pressure necessary under these conditions, the conventional perforated plates are pressed tightly against the adapter end-plate.

DISCLOSURE OF THE INVENTION

It has now been found that a more uniform plug flow and an increase in plate number can be achieved in large-scale chromatography when a liquid gap is provided between the adapter end-plate and the perforated plate during the chromatographic process. This gap is obtained by providing a perforated plate with projections which function to generate a gap, in addition to openings. The positive effect obtained is because liquid that enters the adapter is spread radially through the perforated plate and penetrates the matrix, through the plate openings, in a continuous vertical layer across the end-area of the matrix, resulting in a uniform plug flow and a higher bottom number.

The inventive liquid-chromatographic column is comprised of a tube which contains a chromatographic gel matrix and an adapter

placed at the inlet and the outlet of the column respectively. The invention is characterized in that at least one of the adapters, preferably the inlet adapter, includes a perforated plate and an end-plate, and in that a gap is formed between the
5 end-plate and the perforated plate.

The invention will now be described in more detail with reference to the accompanying drawings, in which

Figure 1 illustrates one embodiment of an inventive column with applied flow;

10 **Figure 2** is a cross-sectional view of the adapter;

Figure 3 is a view of the perforated plate from above; and

Figure 4 illustrates two chromatograms in small and large scale respectively, there being used an inventive column tube in the large-scale test.

15 **Figure 1** illustrates a column tube 1 filled with a matrix 2 and including two adapters 3 and 4 which respectively cover the inlet and outlet areas of the matrix. The adapters 3, 4 include respectively a liquid flow inlet 5 and a liquid flow outlet 6. The flow direction is arrowed in **Figure 1**. Each of the adapters
20 3 and 4 is provided with a conventional sealing element, e.g. O-rings, which seal against the inner surface of the column 1.

The adapters 3 and 4 are shown in larger scale in **Figure 2**, and include an end-plate 7 and a perforated plate 8 provided with openings 8a and projections 8b. The projections may
25 alternatively be disposed on the end-plate of the adapter. The end-plate 7 and the perforated plate 8 define therebetween a gap 9, preferably a gap of from 0.2 to 1.0 mm in depth. The density of the projections is determined by the pressure at which the bed is packed, i.e. the greater the packing pressure,
30 the denser the projections. It is essential that the gap depth will permit the flow to be uniformly distributed through the area of the perforated plate 8 when the column is packed. If the gap depth is excessively small, the flow will be distributed unevenly across the surface area of the plate. The
35 gap depth will normally be > 0.2 mm. The projections serve two purposes. Firstly, the projections will function as gap forming elements, and secondly shall exert the least possible resistance to radial flow, i.e. shall permit the incoming and

outgoing flow to change direction through 90°. Appropriate shapes of the projections are therefore circular, oval, etc. According to one preferred embodiment of the invention, the projections are round.

5 A fine-mesh net 10 is preferably placed against the perforated plate, i.e. against the gel matrix, in a conventional manner not described. The advantage afforded by such a net is that the flow will be spread twice, i.e. once between the end plate and the perforated plate and once between the perforated plate and
10 the net.

The adapter parts are conveniently manufactured separately, and preferably from stainless steel. The parts can be welded together for use in chromatography. The parts can be cleaned very easily and thus prevent undesirable contamination.

15 The inventive column is suitable for matrices having particle sizes of 5-250 μm , particularly for particle sizes of 5-60 μm , and flow rates of 50-1 500 cm/h.

The use of the invention in application will now be illustrated with reference to a number of examples.

20

Example 1

An inventive column tube having an inner diameter of 200 mm and a gel height of 30 mm was packed with 15 μm beads. The
25 column was equalized with 0.5 M NaCl. A trace substance of 2.0 M NaCl with a sample volume corresponding to 1% of the gel volume was introduced to the column, whereafter the column was run with 0.5 M NaCl at a rate of flow of 60 cm/h. The conductivity was registered with a conductivity meter.

30 The peak was shown as a sharp peak and the plate number was calculated as N/L 26 300.

Example 2

This example was run on two scales; a small scale with 2.2 ml
35 gel and a column measuring internally 7.5 mm x 50 mm, and on a large scale with 1570 ml gel and a column having internal dimensions of 200 mm x 50 mm. In the small scale, the column was operated solely with nets, i.e. in the absence of

perforated plates. In the large-scale test there was used a perforated plate in accordance with the invention. The particle size was 30 μm in both cases. The column was equalized with a 20 mM phosphate buffer, pH 6.8. Other conditions:

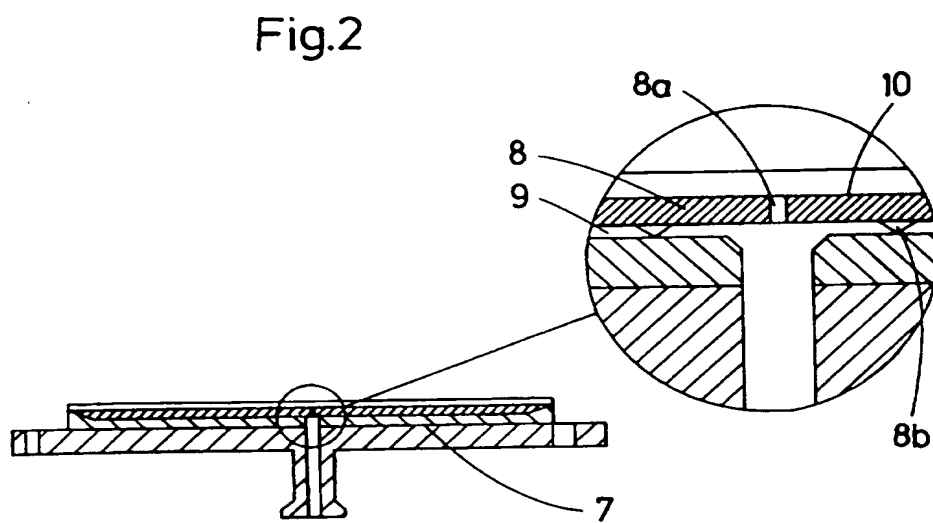
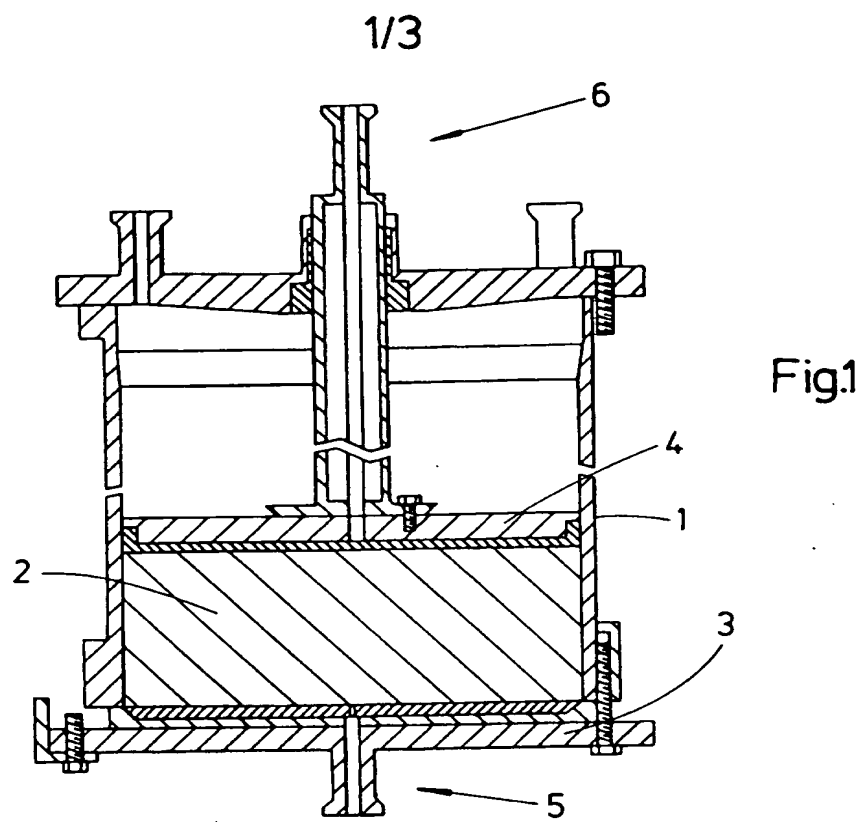
- 5 Sample: ribonuclease A, cytochrome C, lysozyme (3.75:1:1)
 Amount of sample: 0.32 mg/ml gel
 Flow rate: 300 cm/h
 Buffer A: 20 mM phosphate buffer, pH 6.8
 Buffer B: Buffer A + 0.4 M NaCl
10 Gradient: 0-100% B, 10 column volumes.

The results from Example 2 are shown in Figure 4, wherein the upper chromatogram shows the result obtained with the small-scale test, and the lower chromatogram shows the result obtained with the large-scale test. It will be seen that three
15 pronounced peaks occur in both instances. The resolution, i.e. the R_s -value, between the protein peaks for ribonuclease A (peak 1) and cytochrome C (peak 2) is 3.6 in the small scale and 3.6 in the large scale. The resolution between the protein peaks for cytochrome C (peak 2) and lysozyme (peak 3) is 2.2 in
20 the small scale and 2.4 in the large-scale test.

Thus, despite the increase in diameter, the large-scale test gave an equally as good result as and even a better result than the small-scale test. Only unsatisfactory results have been achieved earlier. These results can now be greatly improved in
25 accordance with the invention, with a column provided with adapters that include perforated plates.

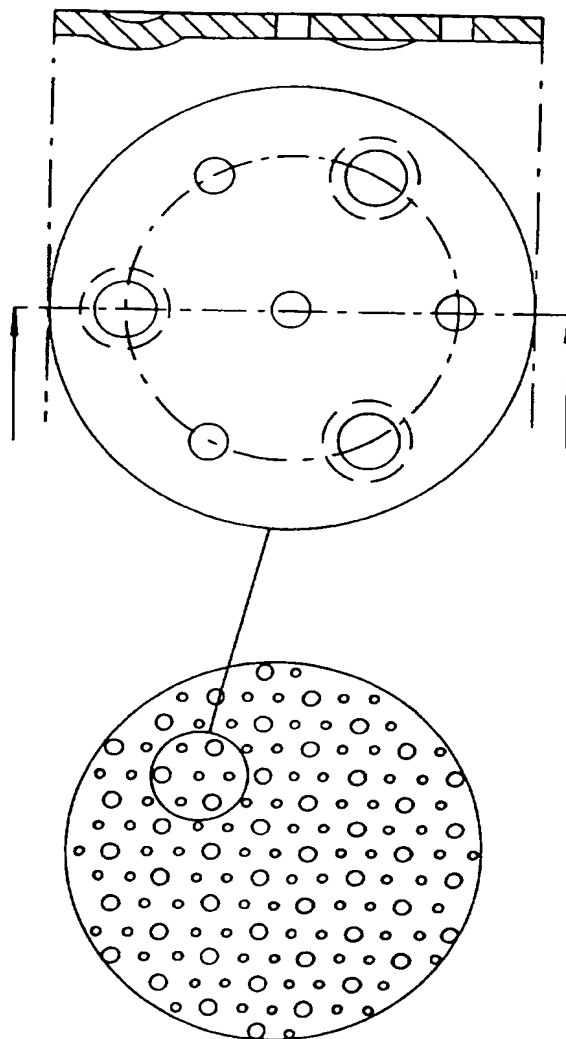
CLAIMS

1. A liquid-chromatography column constructed from a column tube (1) that includes a chromatographic matrix (2), a flow inlet (3) and a flow outlet (4), and an adapter (5 and 6) placed at the inlet and outlet respectively, **characterized** in that respective adapters (5 and 6) include an end-plate (7) and a perforated plate (8) configured to define a gap (9) between the end-plate and the perforated plate (8).
2. A column according to Claim 1, **characterized** in that the gap is created by projections (8b) on the end-plate (7) or on the perforated plate (8).
3. A column according to Claim 1 or 2, **characterized** in that the gap has a depth of 0.2-1.0 mm.
4. A column according to one or more of the preceding claims, **characterized** in that the projections are circular.
5. A column according to one or more of the preceding claims, **characterized** by a net placed between matrix and perforated plate.



2/3

Fig.3



3/3

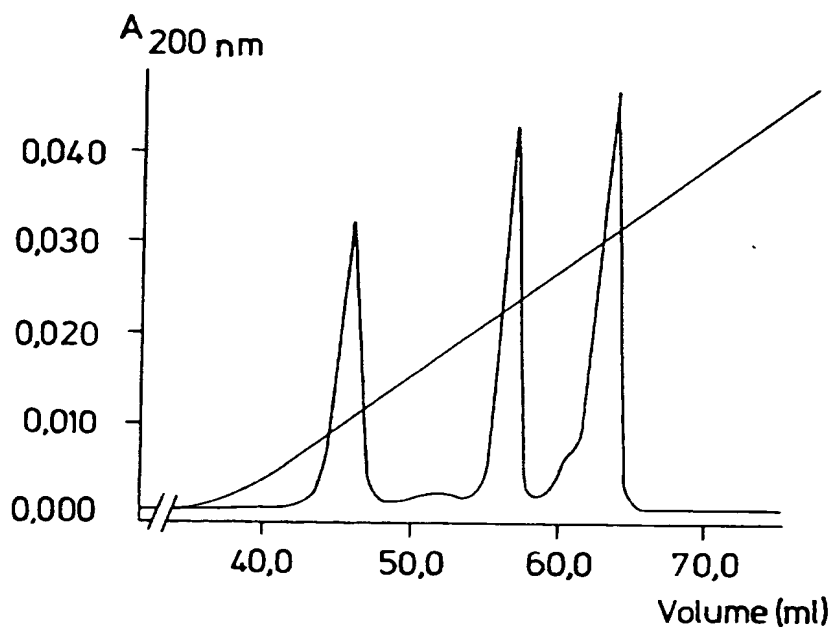


Fig.4a

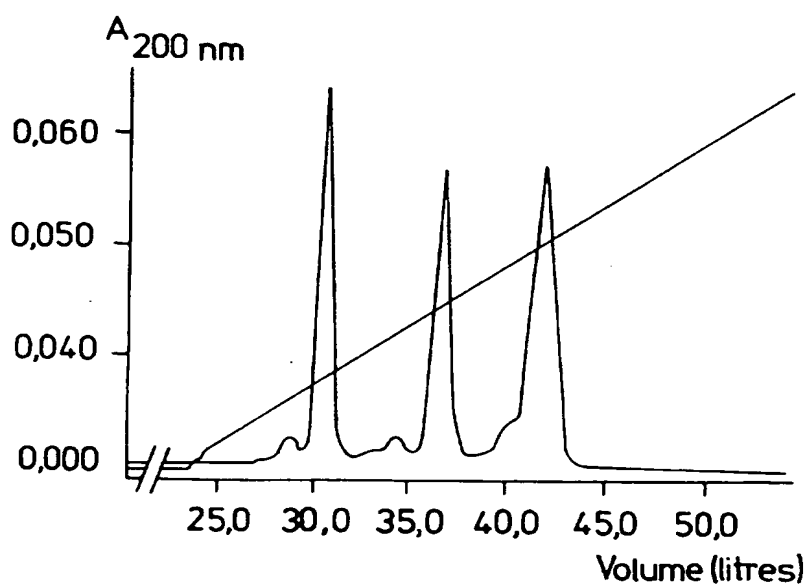


Fig.4b

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 96/00206

A. CLASSIFICATION OF SUBJECT MATTER

IPC6: G01N 30/60, B01D 15/08

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC6: G01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 4470910 A (CLAUDE QUEMERAIS ET AL), 11 Sept 1984 (11.09.84), column 1, line 15 - line 18; column 2, line 11 - line 26; column 5, line 10 - line 27, column 6, line 43 - line 54	1,3,5
Y	--	2
X	US 4557830 A (HATSUKI ONITSUKA ET AL), 10 December 1985 (10.12.85), column 1, line 8 - line 9; column 2, line 55 - line 63; column 5, line 35 - line 53, column 6, line 25 - line 31, column 6, line 50 - line 64	1-3,5
Y	--	2

☒ Further documents are listed in the continuation of Box C.

☒ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

31 May 1996

Date of mailing of the international search report

03 -06- 1996

Name and mailing address of the ISA/

Swedish Patent Office

Box 5055, S-102 42 STOCKHOLM

Facsimile No. +46 8 666 02 86

Authorized officer

Inger Löfgren

Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 96/00206

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 4587014 A (WILLIAM G. AMERICA), 6 May 1986 (06.05.86), figures 1,2, abstract --	1,3,5
X	GB 2248027 A (AMICON LIMITED), 25 March 1992 (25.03.92), page 1, line 9 - line 25; page 4, line 9 - line 16; page 4, line 24 - line 33 -----	1,3,5

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 96/00206

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US-A- 4470910	11/09/84	EP-A,A,B 0089255 FR-A,B- 2522154 JP-A- 58216952	21/09/83 26/08/83 16/12/83
US-A- 4557830	10/12/85	AU-B,B- 546656 AU-A- 2512484 CA-A- 1228531 EP-A,A,A 0123815 JP-C- 1710238 JP-B- 3072338 JP-A- 59199032	12/09/85 01/11/84 27/10/87 07/11/84 11/11/92 18/11/91 12/11/84
US-A- 4587014	06/05/86	CA-A- 1239591 DE-A- 3586068 EP-A,A,A 0161493 JP-A- 60239668	26/07/88 25/06/92 21/11/85 28/11/85
GB-A- 2248027	25/03/92	EP-A,A,A 0476997 JP-A- 4258761 US-A- 5167810	25/03/92 14/09/92 01/12/92